Form PTO-1449 (modified)	Atty. Docket No. IOWA:034US	Serial No. 10/025,187
List of Patents and Publications for Applicant's INFORMATION DISCLOSURE STATEMENT (Use several sheets if necessary)	Applicant Val Sheffield *et al.*	
	Filing Date: December 18, 2001	Group: 1645

U.S. Patent Documents	Foreign Patent Documents	Other Art
See Page 1	*See Page 1*	*See Page 1*

U.S. Patent Documents

Exam. Init.	Ref. Des.	Document Number	Date	Name	Class	Sub Class	Filing Date of App.

Foreign Patent Documents

Exam. Init.	Ref. Des.	Document Number	Date	Country	Class	Sub Class	Translation Yes/No

Other Art (Including Author, Title, Date Pertinent Pages, Etc.)

Exam. Init.	Ref. Des.	Citation
RT	C1	Bruford *et al.*, "Linkage mapping in 29 Bardet-Biedl syndrome families confirms loci in chromosomal regions 11q13, 15q22.3-q23, and 16q21," *Genomics*, 41:93-99, 1997.
	C2	Carmi *et al.*, "Phenotypic differences among patients with Bardet-Biedl syndrome linked to three different chromosome loci," *Am. J. Med. Genet.*, 59:199-203, 1995.
	C3	Carmi *et al.*, "Use of a DNA pooling strategy to identify a human obesity syndrome locus on chromosome 15," *Hum. Mol. Genet.*, 4:9-13, 1995.
	C4	Farag and Teebi, "Bardet-Biedl and Laurence-Moon syndromes in a mixed Arab population," *Clinical Genet.*, 33:78-82, 1988.
	C5	Farag and Teebi, "High incidence of Bardet Biedl syndrome among the Bedouin [letter]," *Clinical Genet.*, 36:463-464, 1989.
/	C6	Katsanis *et al.*, "Mutations in MKKS cause obesity, retinal dystrophy and renal malformations associated with bardet-biedl syndrome," *Nat. Genet.*, 26:67-70, 2000.
	C7	Kwitek-Black *et al.*, "Linkage of Bardet-Biedl syndrome to chromosome 16q and evidence for non-allelic genetic heterogeneity," *Nature Genet.*, 5:392-396, 1993.
	C8	Leppert *et al.*, "Bardet-Biedl syndrome is linked to DNA markers on chromosome 11q and is genetically heterogeneous," *Nature Genet.*, 7:108-112, 1994.
	C9	Nishimura *et al.*, "The forkhead transcription factor gene FKHL7 is responsible for glaucoma phenotypes which map to 6p25," *Nature Genet.*, 19:140-147, 1998.

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EXAMINER: R. Teller	DATE CONSIDERED: 7/14/04

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RT	C10	Sheffield *et al.*, "Identification of a Bardet-Biedl syndrome locus on chromosome 3 and evaluation of an efficient approach to homozygosity mapping," *Hum. Mol. Genet.*, 3(8):1331-1335, 1994.
	C11	Slavotinek *et al.*, "Mutations in MKKS cause bardet-biedl syndrome," *Nature Genet.*, 26:15-16, 2000.
/	C12	Stone *et al.*, "Mutation of a gene encoding a putative chaperonin causes McKusick-Kaufman syndrome," *Nat. Genet.*, 25:79-82, 2000.
	C13	Swiderski *et al.*, "Expression pattern and in situ localization of the mouse homologue of the human MYCO (GLC1A) gene in adult brain," *Mol. Brain Res.*, 68:64-72, 1999.
	C14	Woods *et al.*, "Genetic heterogeneity of Bardet-Biedl syndrome in a distinct Canadian population: evidence for a fifth locus," *Genomics*, 55:2-9, 1999.
	C15	Young *et al.*, "A fifth locus for Bardet-Biedl syndrome maps to chromosome 2q31," *Am. J. Hum. Genet.*, 64:900-904, 1999.

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